REVISED 10 CFR PART 35: MEDICAL USE OF BYPRODUCT MATERIAL

Subpart G: Sealed Sources for Diagnosis

New Subpart G replaces the requirements in the old Subpart H, “Sealed Sources for Diagnosis”

- The following section was deleted:
  - §35.520 Possession of survey instrument
    - §20.1501 requires the licensee to make surveys to demonstrate compliance with Part 20, and requires the licensee to ensure that instruments and equipment used to show compliance with Part 20 are periodically calibrated. In addition, §30.33(a)(2) requires the licensee to have adequate instrumentation.

§35.500 Use of sealed sources for diagnosis

- A licensee shall use only sealed sources for diagnostic uses as approved in the Sealed Source & Device Registry
- Deleted the reference of specific sources and its uses

§35.590 Training for use of sealed sources for diagnosis

- Licensee shall require this AU to be a physician who:
  - 1) Is certified by a medical specialty board whose certification process has been recognized by NRC/Agreement States, or
  - 2) Has completed 8 hours of classroom & laboratory training in basic radionuclide handling techniques specifically applicable to the use of the device